In The Specification:

Please replace the substitute Sequence Listing (1 page) filed on June 13, 2003 with the second substitute Sequence Listing (1 page) filed herewith.

Please replace the paragraph beginning on page 4, line 19 with the following rewritten paragraph:

PCT/EP97/04396 (WO 98/07036) teaches a process for determining the status of an organism by peptide measurement. The reference teaches the measurement of peptides in a sample of the organism which contains both high and low molecular weight peptides and acts indicator of the organism's status. The reference as concentrates on the measurement of low molecular weight peptides, i.e. below 30,000 Daltons, whose distribution serves as representative cross-section of defined controls. Contrary to the methodology of the instant invention, the '396 patent strives to determine the status of a healthy organism, i.e. a "normal" and then use this as reference to differentiate disease states. The present inventors do not attempt to develop a reference "normal", but rather strive to specify particular markers which are evidentiary of at least on specific disease state, whereby the presence of said marker serves as a positive indicator of disease. This leads to a simple method of analysis which can easily be performed by an untrained individual, since there is a positive

correlation of data. On the contrary, the '396 patent requires a complicated analysis by a highly trained individual to determine disease state versus the perception of non-disease or normal physiology.

Please replace the paragraph (first amended on June 13, 2003) beginning at page 19, line 2, with the following rewritten paragraph:

FIGURE 1 is a representation of derived data which characterizes a disease specific marker having a particular sequence (amino acid residues 2-14 of SEQ ID NO: 1) useful in evidencing and categorizing at least one particular disease state. Each patient listed in the data table shows the presence of the disease specific marker (amino acid residues 2-14 of SEQ ID NO: 1) in their serum.

Please replace the paragraph beginning at page 22, line 19, with the following rewritten paragraph:

Chelating Sepharose SEPHAROSE Mini Column

- Dilute Sera in Sample/Running buffer;
- 2. Add Chelating Sepharose SEPHAROSE slurry to column and allow column to pack:
 - 3. Add UF water to the column to aid in packing;

- 4. Add Charging Buffer once water is at the level of the resin surface;
- 5. Add UF water to wash through non bound metal ions once charge buffer washes through;
- 6. Add running buffer to equilibrate column for sample loading;
 - 7. Add diluted serum sample;
 - 8. Add running buffer to wash unbound protein;
- 9. Add elution buffer and collect elution fractions for analysis;
 - 10. Acidify each elution fraction.

Please replace the paragraph (first amended on April 22, 2002 in the Supplemental Preliminary Amendment and again in the Amendment on June 13, 2003) beginning at page 27, line 17 with the following rewritten paragraph:

As a result of these procedures, the disease specific marker consisting of amino acid residues 2-14 of SEQ ID NO:1 was found. This marker is characterized as Alpha Fibrinogen having a molecular weight of about 1350 daltons. The characteristic profile of the marker is set forth in Figure 2. As easily deduced from the data set forth in Figure 1, this marker is indicative of an individual suffering from renal failure or myocardial infarction.

Please replace the paragraph beginning at page 36, line 2, with the following rewritten paragraph:

The instant invention involves the use of a combination of preparatory steps in conjunction with mass spectroscopy and time-of-flight detection procedures to maximize the diversity of biopolymers which are verifiable within a particular sample. The cohort of biopolymers verified within such a sample is then viewed with reference to their ability to evidence at least particular disease state; thereby enabling a diagnostician to gain the ability to characterize either the presence or absence of [said] at least one disease state relative to recognition of the presence and/or the absence of [said] the biopolymer.